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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of: SCHENERMAN, Mark et al.

Art Unit: 1645

Serial No.: 10/751,744

Examiner: Robert A. Zeman

Filed: January 5, 2004

Atty. Docket: AE300US1

For: STABILIZED GLYCOPROTEINS

Confirmation No.: 6583

Mail Stop MISSING PARTS Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

SUBMISSION OF SUBSTITUTE SEQUENCE LISTING INCLUDING STATEMENT per 37 C.F.R. §1.825(c) and (e)

Dear Sir:

Pursuant to the Notice To Comply, mailed May 12, 2006, Applicant hereby provides (1) a copy of the Notice To Comply; (2) a Substitute Paper Copy of the Substitute Sequence Listing; (3) a Computer Readable Form of said Substitute Sequence Listing; (4) an amendment directing entry of the same into the specification. Applicant submits that the amendment is fully supported in the application as filed.

Pursuant to 37 C.F.R. §1.825(c) and (e), the undersigned also states that the Substitute Paper Copy and the Substitute Computer Readable Form are the same and the replacement compact disc includes no new matter.

Respectfully submitted,

Date: May 31, 2006

Janet M. Martineau Attorney for Applicants Reg. No. 46,903

MEDIMMUNE, INC.

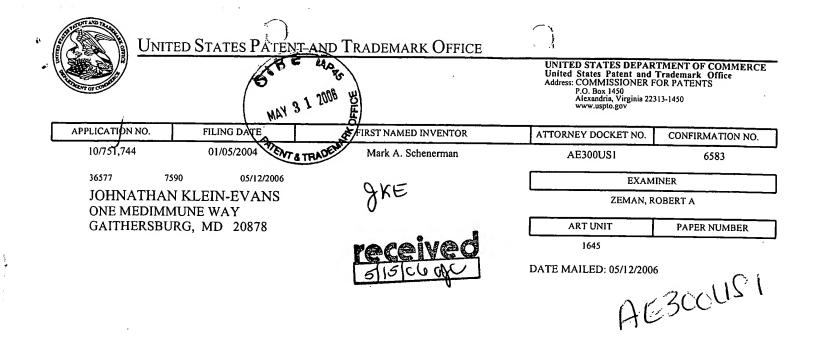
One MedImmune Way Gaithersburg, Maryland 20878 (301) 398-4532 – Tel (301) 398-9306 – Fax

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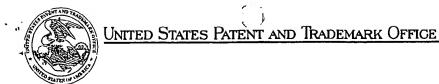
PTO/SB/21 (09-04)

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	Application Number	10/751,744	
TRANSMITTAL	Filing Date	January 5, 2004	
FORM	First Named Inventor	Mark SCHENERMAN	
	Art Unit	1645	
(to be used for all correspondence after initial	filing) Examiner Name	Robert A. Zeman	
Total Number of Pages in This Submission	72 Attomey Docket Number	AE300US1	
ENCLOSURES (Check all that apply)			
Fee Transmittal Form	Drawing(s)	After Allowance Communication to TC	
Fee Attached	Licensing-related Papers	Appeal Communication to Board of Appeals and Interferences	
Amendment/Reply After Final Affidavits/declaration(s) Extension of Time Request Express Abandonment Request Information Disclosure Statement Certified Copy of Priority Document(s) Reply to Missing Parts/ Incomplete Application Reply to Missing Parts under 37 CFR 1.52 or 1.53	Petition Petition to Convert to a Provisional Application Power of Attorney, Revocation Change of Correspondence Ar Terminal Disclaimer Request for Refund CD, Number of CD(s) Landscape Table on CD Remarks	Appeal Communication to TC (Appeal Notice, Brief, Reply Brief) Proprietary Information	
SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT			
MEDIMMUNE, INC.			
Signature DM 1	ul i		
Printed name Variet MARTINEAU			
Date May 31, 2006	R	eg. No. 46,903	
CERTIFICATE OF TRANSMISSION/MAILING			
I hereby certify that this correspondence is being facsimile transmitted to the USPTO or deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450 on the date shown below:			
Signature			
Typed or printed name		Date	

This collection of information is required by 37 CFR 1.5. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and1.14. This collection is estimated to 2 hours to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.



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APPLICATION NO. CONTROL NO. 10/751,744

FIRST NAMED INVENTOR /PATENT IN REEXAMINATION ATTORNEY DOCKET NO.

EXAMINER

Robert A. Zeman

ART UNIT PAPER

1645

DATE MAILED:

Please find below and/or attached an Office communication concerning this application or proceeding.



Commissioner of Patents

This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 C.F.R. § 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 C.F.R. §§ 1.821-1.825 for the reason(s) set forth on the attached Notice To Comply With Requirements For Patent Applications Containing Nucleotide Sequence And/Or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be examined under 35 U.S.C. §§ 131 and 132.

APPLICANT IS GIVEN ONE MONTH FROM THE DATE OF THIS LETTER WITHIN WHICH TO COMPLY WITH THE SEQUENCE RULES, 37 C.F.R.. §§ 1.821-1.825. Failure to comply with these requirements will result in ABANDONMENT of the application under 37 C.F.R. § 1.821(g). Extensions of time may be obtained by filing a petition accompanied by the extension fee under the provisions of 37 C.F.R. § 1.136. In no case may an applicant extend the period for response beyond the six month statutory period. Direct the response to the undersigned. Applicant is requested to return a copy of the attached Notice to Comply with the response.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Robert A. Zeman whose telephone number is (571) 272-0866.

ROBERT ZEMAN PATENT EXAMINER

(iii) plication No.: 10/751,744

NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

wing	reason(s):		
X X	1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).		
	2. This application does not contain, as a separate part of the disclosure on paper copy, a □Sequence Listing□ as required by 37 C.F.R. 1.821(c).		
	3. A copy of the Sequence Listing in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).		
	4. A copy of the Sequence Listing in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up Raw Sequence Listing.		
	5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).		
	6. The paper copy of the Sequence Listing is not the same as the computer readable from of the □Sequence Listing□ as required by 37 C.F.R. 1.821(e).		
X	7. Other: the specification contains sequences without the proper sequence identifiers (see page 81 for example.		
Applicant Must Provide:			
X	An initial or <u>substitute</u> computer readable form (CRF) copy of the Sequence Listing		
X	An initial or <u>substitute</u> paper copy of the Sequence Listing, as well as an amendment directing its entry into the specification.		
X	A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).		
Fo	r questions regarding compliance to these requirements, please contact:		
	For Rules Interpretation, call (703) 308-4216		
For CRF Submission Help, call (703) 308-4212			
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